AMENDMENTS TO THE CLAIMS

Claims 1-16 (Previously cancelled)

Claim 17. (Currently amended) A compound having the structure:

$$H \xrightarrow{A^{1} A^{2}} H$$

$$H \xrightarrow{N \quad H} H$$

or an optical isomer, diastereomer, enantiomer, or pharmaceutically acceptable salt, or amide, ester, or imide susceptible to being cleaved *in vivo* by a mammalian subject to yield the compound, wherein:

(a) A^1 and A^2 are each, independently, selected from the group consisting of a hydrogen atom and a group having the structure:

$$\begin{cases} \begin{pmatrix} R^1 \\ C \\ R^1 \end{pmatrix} D^1 - D^2 - R^2 \\ x \end{cases}$$

with the proviso that at A¹ and A² are not both hydrogen atoms, and wherein:

- (i) each R¹ is independently selected from the group consisting of a hydrogen atom and a hydroxyl group;
- (ii) x is 0 or 1;
- (iii) each R² is independently selected from the group consisting of:

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$$\begin{array}{c|c} & & & & \\ & &$$

wherein:

- (a) a is at least 2;
- (b) b is at least 2;
- (c) c is 1 to 3;
- (d) d is 1 to 3; and
- (e) R¹² and R¹³ are each independently selected from the group consisting of hydrocarbon groups and substituted hydrocarbon groups; and
- (iv) D^1 -and D^2 -are each independently selected from the group consisting of C(O)- and NH; with the proviso that wherein when D^1 is NH- then D^2 is C(O); and wherein when D^2 is NH- then D^1 is C(O); and,
- (v) D^2 is -NH-.
- (b) A^3 has the structure:

$$\begin{array}{c} \begin{array}{c} \\ \\ \end{array} \begin{array}{c} \\ \end{array} \begin{array}{c} \\ \end{array} \begin{array}{c} \\ \\ \end{array} \begin{array}{c} \\ \\ \end{array} \begin{array}{c} \\ \end{array} \begin{array}{c} \\ \\ \end{array} \begin{array}{c} \\ \end{array} \begin{array}{c} \\ \\ \end{array} \begin{array}{c} \\ \\ \end{array} \begin{array}{c} \\ \end{array} \begin{array}{c} \\ \\ \end{array} \begin{array}{c} \\ \end{array} \begin{array}{c} \\ \end{array} \begin{array}{c} \\ \\ \end{array} \begin{array}{c} \\$$

wherein:

- (i) each R¹ is independently selected from the group consisting of a hydrogen atom and a hydroxyl group;
- (ii) t is from 0 to 6;

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(iii)
$$D^4$$
 is $-CH(R^1)$ -;
(iv) D^5 is $-OR^6$; and

- (v) R⁶ is selected from the group consisting of a carbocyclic group, a substituted carbocyclic group, an aromatic group, and a substituted aromatic group.
- Claim 18. (Previously added) The compound according to Claim 17 wherein x is 1.
- Claim 19. (Previously added) The compound according to claim 17 wherein x is 0. Claim 20. (Currently cancelled)
- Claim 21. (Currently cancelled)
- Claim 22. (Currently cancelled)
- Claim 23. (Previously amended) The compound according to Claim 17 wherein t is 0 to 2.
- Claim 24. (*Previously added*) The compound according to Claim 17 wherein R⁶ is a substituted aromatic group.
- Claim 25. (Previously added) A composition comprising:
 - (a) the compound according to Claim 1; and
 - (b) a pharmaceutically acceptable carrier.

Claim 26. (*Previously added*) A method selected from the group consisting of treating multidrug resistance, inhibiting transport protein activity, combinations thereof, comprising administering to a mammal in need of such treatment or inhibition an effective amount of the composition according to Claim 2.